PRODUCT MONOGRAPH

INCLUDING PATIENT MEDICATION INFORMATION

PrTYRUKO

Natalizumab for injection

Concentrate for solution for intravenous infusion

300 mg/15 mL

Monoclonal antibody

Polpharma Biologics S.A.
Ul. Trzy Lipy 3
Gdansk
Pomorskie
80-172
Poland

Imported and distributed by:
Sandoz Canada Inc.
4600, rue Armand-Frappier
Saint-Hubert, Quebec J3Z 1G5

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Recent Major Label Changes

Not applicable	

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Sections or subsections that are not applicable at the time of authorization are not listed.

Part I: Health Professional Information

TYRUKO (natalizumab for injection) is a biosimilar biologic drug (biosimilar) to TYSABRI® (natalizumab for injection). A biosimilar is a biologic drug that was granted authorization based on a demonstration of similarity to a version previously authorized in Canada, known as the reference biologic drug.

1 Indications

TYRUKO (natalizumab for injection) is indicated:

- as monotherapy (i.e. single disease-modifying agent) for the treatment of patients with the relapsing-remitting form of multiple sclerosis (MS) to reduce the frequency of clinical exacerbations, to decrease the number and volume of active brain lesions identified on magnetic resonance imaging (MRI) scans and to delay the progression of physical disability. TYRUKO is generally recommended in MS patients who have had an inadequate response to, or are unable to tolerate, other therapies for multiple sclerosis.

Safety and efficacy in patients with chronic progressive multiple sclerosis, and in geriatric and pediatric patients, have not been established.

The efficacy of natalizumab for a treatment duration beyond 2 years has not been determined.

TYRUKO should be used by physicians who have sufficient knowledge of multiple sclerosis and who have familiarized themselves with the efficacy/safety profile of TYRUKO.

TYRUKO is only available through a controlled distribution program called Sandoz PLUS Support Program. Under this program, only prescribers and pharmacies registered with the program are able to prescribe and dispense the product. In addition, TYRUKO can only be dispensed to patients who are registered and meet all the conditions of the Sandoz PLUS Support Program. Please call 1-888-449-7673 or log onto website http://www.my-sandoz-plus.ca.

1.1 Pediatrics

Pediatrics (<18 years of age): Safety and effectiveness of natalizumab in pediatric patients with multiple sclerosis have not been established. Therefore, Health Canada has not authorized an indication for pediatric use.

1.2 Geriatrics

Geriatrics (>65 years of age): Clinical studies of natalizumab did not include sufficient numbers of patients aged 65 years and over to determine whether they respond differently than younger patients.

2 Contraindications

- TYRUKO is contraindicated in patients who are hypersensitive to this drug or to any ingredient in the formulation, including any non-medicinal ingredient, or component of the container. For a complete listing, see 6 Dosage Forms, Strengths, Composition And Packaging.
- TYRUKO is contraindicated in patients who have or have had progressive multifocal leukoencephalopathy (PML). See 7 Warnings and Precautions.

- TYRUKO is contraindicated in patients who are immunocompromised, including those immunocompromised due to immunosuppressant or antineoplastic therapies, or immunodeficiencies (HIV, leukemias, lymphomas, etc.). See 7 Warnings and Precautions.

3 Serious Warnings and Precautions Box

Serious Warnings and Precautions

- **Treatment with TYRUKO (natalizumab for injection) has been associated with an increased risk of progressive multifocal leukoencephalopathy (PML) and/or granule cell neuronopathy (GCN) secondary to opportunistic infection caused by John Cunningham virus (JC virus). Both PML and JCV GCN can cause disability or death. See 2 Contraindications, 7 Warnings and Precautions, Infections, and 8 Adverse Reactions.**
- **Healthcare professionals should monitor patients on TYRUKO for any new sign or symptom that may be suggestive of PML and/or JCV GCN. TYRUKO dosing should be withheld immediately at the first sign or symptom suggestive of either PML or JCV GCN. Diagnosis and management of GCN should follow guidance provided for PML.**

4 Dosage and Administration

4.1 Dosing Considerations

TYRUKO (natalizumab for injection) should be administered by a healthcare professional.

- Patients should be observed during the infusion and for 1 hour after the infusion is complete for signs and symptoms of infusion reactions. Promptly discontinue the infusion upon the first observation of any signs or symptoms consistent with a hypersensitivity reaction (see 7 Warnings and Precautions, Immune, Hypersensitivity Reactions).
- Dilute only with 0.9% Sodium Chloride Injection, USP.
- TYRUKO contains 52 mg sodium per vial. When diluted in 100 mL of 0.9% Sodium Chloride Injection, USP, TYRUKO contains 406 mg sodium per dose. This should be taken into consideration by patients on a controlled sodium diet.

4.2 Recommended Dose and Dosage Adjustment

The recommended dose of TYRUKO is 300 mg IV infusion every 4 weeks. Do not administer TYRUKO as an IV push or bolus injection.

Health Canada has not authorized an indication for pediatric use (see 1.1 Pediatrics).

Pharmacokinetics of natalizumab in patients with renal or hepatic insufficiency have not been studied.

4.3 Reconstitution

Parenteral Products:

Table 1 – Reconstitution

Vial Size	Volume of Diluent	Approximate Volume for infusion	Concentration per mL
15 mL	100 mL 0.9% Sodium Chloride Injection, USP	115 mL	2.6 mg

Intravenous infusion: If not used immediately, store the TYRUKO solution for infusion at 2°C - 8°C. TYRUKO solution for infusion must be administered within 24 hours of preparation.
DO NOT FREEZE. See 11 Storage, Stability and Disposal.

4.4 Administration

Dilution:

Parenteral Products:

Use aseptic technique when preparing TYRUKO solution for IV infusion. Each vial contains a single dose and is intended for single patient use only.

TYRUKO is a clear, colourless to slightly cloudy liquid. Inspect the TYRUKO vial for particulate material prior to dilution and administration. If visible particulates are observed and/or the liquid in the vial is discoloured, the vial must not be used. Do not use TYRUKO beyond the expiration date on the carton or vial.

To prepare the solution, withdraw 15 mL of TYRUKO concentrate from the vial using a sterile needle and syringe. Inject the concentrate into 100 mL 0.9% Sodium Chloride Injection, USP. No other IV diluents may be used to prepare the TYRUKO solution.

Gently invert the TYRUKO solution to mix completely. Do not shake. Inspect for particulate material prior to administration.

Infuse over approximately 1 hour. Observe patients during the infusion and for 1 hour after the infusion is completed for signs and symptoms of infusion reactions. After the first 12 TYRUKO doses, patients should continue to be observed during infusion. If patients have not experienced any infusion reactions including hypersensitivity, the post dose observation time may be adjusted for the 13th and subsequent infusions according to clinical judgement.

After the infusion is complete, flush with 0.9% Sodium Chloride Injection, USP. Other medications should not be injected into infusion set side ports or mixed with TYRUKO.

4.5 Missed Dose

If a dose of TYRUKO is missed, TYRUKO should be administered as soon as possible, and dosing should continue at the prescribed frequency.

5 Overdosage

Safety of doses higher than 300 mg has not been adequately evaluated. The maximum amount

of natalizumab that can be safely administered has not been determined.

For the most recent information in the management of a suspected drug overdose, contact your regional poison control centre or Health Canada's toll-free number, 1-844 POISON-X (1-844-764-7669).

6 Dosage Forms, Strengths, Composition And Packaging

To help ensure the traceability of biologic products, including biosimilars, health professionals should recognise the importance of recording both the brand name and the non-proprietary (active ingredient) name as well as other product-specific identifiers such as the Drug Identification Number (DIN) and the batch/lot number of the product supplied.

Table 2 – Dosage Forms, Strengths, Composition and Packaging

Route of Administration	Dosage Form / Strength/Composition	Non-medicinal Ingredients
Intravenous infusion	Concentrate for solution / 300 mg per 15 mL	L-histidine, L-histidine hydrochloride monohydrate, polysorbate 80, sodium chloride, water for injection

TYRUKO (natalizumab for injection) concentrate is supplied as 300 mg natalizumab in a sterile, single-use vial free of preservatives.

Each package contains one vial.

Each 15 mL dose also contains (pH 5.7):

131.49 mg sodium chloride, USP/ Ph.Eur

6.36 mg L-histidine, USP/ Ph.Eur

22.86 mg L-histidine hydrochloride, monohydrate/ Ph.Eur

3.0 mg polysorbate 80, USP/ Ph.Eur

Water for Injection, USP/ Ph.Eur

7 Warnings and Precautions

Please see 3 Serious Warnings And Precautions Box.

General

Before initiation of treatment with TYRUKO (natalizumab for injection), a recent magnetic resonance image (MRI) should be available. This MRI may be helpful in differentiating subsequent MS symptoms from PML. For diagnosis of PML, an evaluation that includes a magnetic resonance imaging (MRI) scan of the brain and, when indicated, cerebrospinal fluid analysis for JC viral DNA are recommended. See 7 Warnings and Precautions, Infections).

Patients who are prescribed TYRUKO are to be enrolled in the Sandoz PLUS Support Program –

a registry of Canadian patients. To enroll, physicians fax an enrollment form to 1-888-449-7673. Sandoz PLUS Support Program is a comprehensive program associated with the prescribing, administration and monitoring of patients who receive TYRUKO treatment.

Educational Guidance: Prescribers and healthcare professionals are educated regarding the appropriate use of TYRUKO, and information is updated regularly. Prescribers are expected to educate patients on the benefits and risks of treatment, especially the risk of PML. Patients should remain under medical supervision while receiving TYRUKO and should be evaluated by the prescriber every six months.

At 24 months of treatment, physicians should inform patients again about the risks of TYRUKO, including that the risk of PML increases with longer treatment duration. Physicians should obtain consent from their patients for continuation of treatment. At 24 months of treatment, the patient, their partner and/or caregiver should be re-instructed about the early signs and symptoms of PML.

Carcinogenesis and Mutagenesis

No clastogenic or mutagenic effects of natalizumab were observed in the Ames human chromosomal aberration assays. Natalizumab showed no effects on in vitro assays of α4-integrin-positive human tumour line proliferation/cytotoxicity. Xenograft transplantation models in SCID and nude mice with two α4-integrin-positive human tumour lines (leukemia, melanoma) demonstrated no increase in tumour growth rates or metastasis resulting from natalizumab treatment (See 16 Non-Clinical Toxicology).

No differences in incidence rates or the nature of malignancies between natalizumab and placebo- treated patients were observed over 2 years of treatment. However, observation over longer treatment periods is required before any effect of natalizumab on malignancies can be excluded. Should a malignancy develop, TYRUKO therapy should be withheld at least until appropriate treatment has been initiated for the malignancy and the benefit and risks of resuming TYRUKO therapy have been deemed to be acceptable by the treating physician.

Driving and Operating Machinery

No studies on the effects on the ability to drive and use machines have been performed with natalizumab. However, given that dizziness has been very commonly reported, patients who experience this adverse reaction should be advised not to drive or use machines until it has resolved.

Hematologic

In clinical trials, natalizumab was observed to induce increases in circulating lymphocytes, monocytes, eosinophils and nucleated red blood cells. During phase 3 clinical trials, cell counts were measured every 12 weeks. The largest cell increases were seen in lymphocytes, which were found to be elevated within 12 weeks after initiating natalizumab treatment, reaching a plateau by 24 weeks. Although elevated, mean cell counts remained within the normal range. Observed increases persisted during natalizumab exposure, but were reversible, returning to baseline levels usually within 16 weeks after the last dose. Elevations of neutrophils were not observed. Natalizumab may induce mild decreases in hemoglobin levels (mean decrease of 6.0 g/L) that are frequently transient. Hemoglobin levels returned to pretreatment values,

usually within 16 weeks of last dose of natalizumab and the changes were not associated with clinical symptoms.

In post-marketing experience, there have been reports of uncommon frequency of thrombocytopenia and immune thrombocytopenic purpura (ITP), some of which were serious. Symptoms of thrombocytopenia may include spontaneous bruising, abnormal bleeding, and petechiae. Delay in the diagnosis and treatment of thrombocytopenia may lead to serious and life-threatening sequelae. If thrombocytopenia is identified, discontinuation of natalizumab should be considered.

Hepatic

In post-marketing experience, there have been rare reports of clinically significant liver injury, including markedly elevated serum hepatic enzymes and elevated total bilirubin, as early as 6 days after the first dose; signs of liver injury have also been reported for the first time after multiple doses. In some patients, liver injury recurred upon rechallenge. Some cases occurred in patients with pre-existing liver disease or in the presence of other drugs that have been associated with hepatic injury. Patients with history of liver disease, alcohol abuse, and/or treatment with other therapies that are known to cause liver injury should be carefully evaluated prior to commencement of treatment with natalizumab and closely monitored for possible liver damage during and after treatment. The combination of transaminase elevations and elevated bilirubin without evidence of obstruction is generally recognized as an important predictor of severe liver injury that may lead to death or the need for a liver transplant in some patients.

TYRUKO should be discontinued in patients who develop jaundice or other evidence of clinically significant liver injury, e.g., 5-fold or greater elevation in serum hepatic enzymes, and the patient should be fully evaluated. In cases with no other identifiable cause, TYRUKO should be permanently discontinued.

Immune

Hypersensitivity Reactions:

Natalizumab has been associated with hypersensitivity reactions, which occurred at an incidence of 4%, including serious systemic reactions (e.g. anaphylaxis), which occurred at an incidence of <1%. These reactions usually occurred within 2 hours of the start of the infusion. Symptoms associated with these reactions included urticaria, dizziness, fever, rash, rigors, pruritus, nausea, flushing, hypotension, dyspnea and chest pain. Generally, these reactions are associated with antibodies to natalizumab. The risk for hypersensitivity was greatest with early infusions and in patients re-exposed to natalizumab following an initial short exposure (up to three infusions) and extended period (three months or more) without treatment. If a hypersensitivity reaction occurs, discontinue administration of natalizumab immediately and initiate appropriate therapy.

Physicians should inform patients about the importance of uninterrupted dosing, particularly in the early months of treatment.

Post-Marketing Experience: There have been reports of hypersensitivity reactions which have been associated with one or more of the following: hypotension, hypertension, chest pain,

chest discomfort, and dyspnea.

Immunosuppression:

The safety and efficacy of natalizumab in combination with antineoplastic or immunosuppressive agents have not been established. Concurrent use of these agents with natalizumab may increase the risk of infections, including opportunistic infections, over the risk observed with use of natalizumab alone. The risk of PML is also increased in patients who have been treated with an immunosuppressant prior to receiving natalizumab (see Contraindications).

No studies have been performed to evaluate the efficacy and safety of natalizumab when switching patients from disease-modifying therapies with an immunosuppressant effect to natalizumab. It is unknown if patients switching from these therapies to natalizumab have an increased risk of PML. Therefore, these patients should be monitored more frequently, similarly to patients switching from immunosuppressants to natalizumab (see Warnings and Precautions, Infections, Magnetic Resonance Imaging Screening for PML).

In clinical studies for conditions other than MS, opportunistic infections (*e.g. pneumocystis carinii pneumonia, pulmonary mycobacterium avium intracellulare, bronchopulmonary aspergillosis and burkholderia cepacia*) have been uncommonly observed in patients receiving natalizumab; some of these patients were receiving concurrent immunosuppressants (see Adverse Reactions). In pivotal clinical trials (1801 and 1802), concomitant treatment of relapses with a short course of corticosteroids was not associated with an increased rate of infection in patients treated with natalizumab as compared with placebo.

Immunizations:

In a randomized, open label study of 60 patients with relapsing MS for tetanus antibody response, post-immunization antibody levels for natalizumab-treated patients were lower compared to controls (analysis based on 40/60 (67%) subjects available for the primary analysis), and a slower and reduced humoral immune response to neoantigen (keyhole limpet haemocyanin) was observed (analysis based on 42/60 (70%) subjects available for the primary analysis). The clinical significance of this is unknown. Live vaccines have not been studied.

Infections

Progressive Multifocal Leukoencephalopathy:

Use of natalizumab has been associated with an increased risk of progressive multifocal leukoencephalopathy (PML), an opportunistic infection caused by John Cunningham Virus (JC virus). PML can cause severe disability or death.

Risk of PML

Three factors that are known to increase the risk of PML in natalizumab-treated patients have been identified:

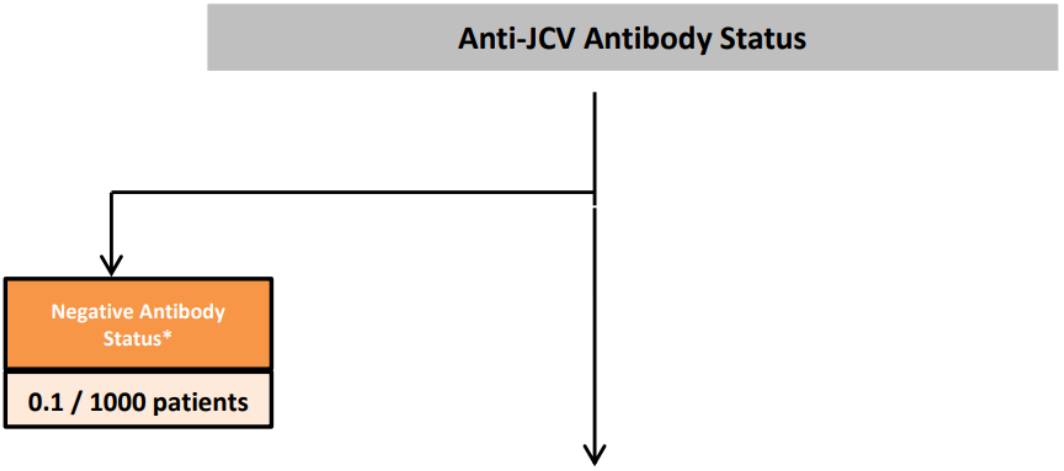
- The presence of anti-JCV antibodies. Testing for the presence of anti-JCV antibodies should only be performed using a validated JCV assay test (see Testing for Anti-JCV Antibody);
- Longer treatment duration, especially beyond 24 months; and

- Prior treatment with an immunosuppressant (IS), which appears independent of natalizumab treatment duration.

The PML Risk Estimates Algorithm (Figure 1) summarizes PML risk by anti-JCV antibody status, prior IS use and duration of treatment (by year of treatment) and stratifies this risk by anti-JCV antibody level (index value), as derived from a validated JCV assay test (see Testing for Anti-JCV Antibody). The risk estimates are forward-looking in yearly intervals (e.g., the risk estimate corresponding to the 25-36 month natalizumab exposure period is the PML risk estimated for the next year for patients treated for 24 months with natalizumab). The individual treatment length of each patient is taken into consideration accounted for dropouts (e.g., treatment discontinuations).

Patients who are anti-JCV antibody negative are at a significantly lower risk of developing PML. Patients who have all three factors (anti-JCV antibody positive **and** prior IS use **and** duration of TYRUKO treatment >2 years) are at higher risk of PML. In patients not previously treated with IS, the level of anti-JCV antibodies (index value) can further stratify risk for PML. Index values equal to or below 0.9 are associated with a PML incidence of less than 1 per 1000 patients; PML risk increases substantially at index values above 1.5. The risks and benefits of continuing treatment with TYRUKO should be carefully considered in patients who have all three risk factors for PML or who have no prior IS use and have an index value of greater than 1.5 and more than 2 years of treatment with TYRUKO.

Figure 1: PML Risk Estimates Algorithm



Natalizumab Exposure	PML risk estimates per 1000 patients				
	Patients without prior IS use ***				Patients with prior IS use
	No index value	Antibody Index ≤ 0.9	Antibody Index > 0.9 ≤ 1.5	Antibody Index > 1.5	
1-12 months	0.1	0.1	0.1	0.2	0.3
13-24 months	0.6	0.1	0.3	0.9	0.4
25-36 months	2	0.2	0.8	3	4
37-48 months	4	0.4	2	7	8
49-60 months	5	0.5	2	8	8
61-72 months	6	0.6	3	10	6

*The risk of PML in anti-JCV antibody negative patients were estimated based on post marketing data from approximately 125,000 natalizumab exposed patients.
**PML risk estimates in anti-JCV antibody positive patients were based on the pooled cohort of 21,696 patients who participated in 3 observational studies (STRATIFY-2, TOP and TYGRIS) and 1 clinical study (STRATA).
***The majority of the prior IS use from these studies included the following 5 IS therapies: mitoxantrone, methotrexate, azathioprine, cyclophosphamide and mycophenolate.

The estimates of PML risk as derived from clinical and observational trial data have been consistent with postmarketing data.

Testing for Anti-JCV Antibody

Infection by the JC virus is required for the development of PML. Anti-JCV antibody negative status indicates that exposure to the JC virus has not been detected; such patients are still at risk for the development of PML due to the potential for a new JCV infection, fluctuating antibody status or a false negative test result. For purposes of risk assessment, a patient with a positive anti-JCV antibody test at any time is considered anti-JCV antibody positive regardless of the results of any prior or subsequent anti-JCV antibody testing.

Testing for serum anti-JCV antibody status (using a validated JCV assay test; see details below) prior to initiating TYRUKO therapy or in patients receiving TYRUKO with an unknown antibody

status is recommended. In addition, anti-JCV antibody retesting is recommended for patients with anti- JCV antibody negative status and for those anti-JCV antibody positive patients with lower index value, since the antibody status or index value may change. Retesting of patients who are anti- JCV antibody negative, every 6 months, is recommended. Patients with lower index values who have not had prior IS use should be retested periodically (e.g., every 6 months). Anti-JCV antibody negative patients may still be at risk of PML for reasons such as new JCV infection, fluctuating antibody status or a false negative test result. Based on a postmarketing study examining longitudinal antibody status over 18 months, there was approximately an 11% annual change in serostatus from anti-JCV antibody negative to positive. Testing should only be performed using an ELISA assay that has been validated for use in MS patients.

Please login using this link: https://immunowell-jcv-portal.com for details of the anti-JCV antibody testing service to patients considering or on Tyruko treatments.

The anti-JCV antibody assay (ELISA) should not be used to diagnose PML. Use of plasmapheresis (PLEX) or intravenous immunoglobulin (IVIg) can affect meaningful interpretation of serum anti-JCV antibody testing. Anti-JCV antibody testing should not be performed during or for at least two weeks following PLEX due to the removal of antibodies from the serum or within 6 months of IVIg (i.e., 6 months = 5 x half-life for immunoglobulins).

Magnetic Resonance Imaging Screening for PML

Before initiation of treatment with TYRUKO, a recent magnetic resonance imaging (MRI) scan should be available. Pre-treatment investigations (e.g., MRI) may be helpful in the evaluation of patients who develop signs or symptoms suggestive of PML. More frequent monitoring (e.g., every 3-6 months) should be considered for patients at higher risk of PML. This includes:

- Patients who have all three risk factors for PML (i.e., are anti-JCV antibody positive **and** have received more than 2 years of TYRUKO therapy, **and** have received prior immunosuppressant therapy),

Or

- Patients with an anti-JCV antibody index value of greater than 1.5 without prior history of immunosuppressant therapy and more than 2 years of natalizumab treatment.

PML in the absence of symptoms can be detected on MRI and must be confirmed by presence of JCV DNA in CSF or brain biopsy.

Assessing for PML

Typical symptoms of PML are diverse, progress over days to weeks, and include progressive weakness on one side of the body or clumsiness of limbs, disturbance of vision, and changes in thinking, memory and orientation leading to confusion and personality changes. **Withhold TYRUKO dosing immediately at the first sign or symptom suggestive of PML.**

Healthcare professionals should be particularly alert to symptoms suggestive of PML that the patient may not notice (e.g. cognitive or psychiatric symptoms). Patients should also be advised to inform their partner or caregivers about their treatment, since they may notice

symptoms that the patient doesn't.

Monitoring for PML should continue while the patient is receiving TYRUKO therapy and for a period of 6 months following treatment. PML has been reported following discontinuation of TYRUKO in patients who did not have findings suggestive of PML at the time of discontinuation. Patients and healthcare professionals should continue to be vigilant for any new signs or symptoms that may be suggested of PML for approximately 6 months following discontinuation of TYRUKO (see 7 Warnings and Precautions, Stopping TYRUKO Therapy).

At 2 years of treatment, physicians should inform patients again about the risks of TYRUKO, including that the risk of PML increases with longer treatment duration. Physicians should obtain consent from their patients for continuation of treatment. At 2 years of treatment, the patient, their partner and/or caregiver should be re-instructed about the early signs and symptoms of PML.

TYRUKO should be suspended immediately at the first signs or symptoms suggestive of PML and an evaluation that includes a magnetic resonance imaging (MRI) scan of the brain should be performed. When indicated, cerebrospinal fluid analysis for JC viral DNA is recommended to confirm a diagnosis of PML. If initial investigations prove negative but clinical suspicion for PML still remains, TYRUKO should not be restarted and repeat investigations should be undertaken.

Patients being treated with TYRUKO should be instructed to report any new neurological signs or symptoms to their physician.

There are no known interventions that can reliably prevent PML or adequately treat PML if it occurs. It is not known whether early detection of PML and discontinuation of TYRUKO will mitigate the disease. In the post-marketing setting, early detection of PML and suspension of natalizumab therapy may have contributed to improved survival rates from PML compared with the natalizumab pre-approval clinical trial PML cases. Plasma exchange has been used to reduce the serum levels of natalizumab. (See PML and IRIS (Immune Reconstitution Inflammatory Syndrome) below.)

It is unclear whether the risk of PML is increased in MS patients treated with natalizumab in combination with interferon beta compared to natalizumab alone. TYRUKO should not be used in combination with other immunosuppressive or immunomodulatory agents, regardless of their class.

PML and IRIS (Immune Reconstitution Inflammatory Syndrome)

In natalizumab treated patients who developed PML, Immune Reconstitution Inflammatory Syndrome (IRIS) has been described following discontinuation or removal of natalizumab (by plasma exchange); this can lead to serious neurological complications. In patients who have undergone plasma exchange, IRIS has occurred within days to several weeks. IRIS presents as a worsening in neurological status that may be rapid, as a result of the sudden reconstitution of immune function. It can lead to serious neurological complications and may be fatal.

Monitoring for development of IRIS and appropriate treatment of the associated inflammatory reaction involving the brain should be undertaken.

JCV Granule Cell Neuronopathy (GCN):

JCV also causes granule cell neuronopathy (GCN) which has been reported in patients treated with natalizumab. Symptoms of JCV GCN include progressive unsteadiness in walking (ataxia), slurred slow speech with loss of normal rhythm (dysarthria), and/or incoordination of movements, and diagnosis and management of JCV GCN should follow guidance provided for PML (see Adverse Reactions, Post Market Adverse Drug Reactions, Infections).

Herpes Infections:

Natalizumab increases the risk of developing encephalitis and meningitis caused by herpes simplex and varicella zoster viruses. Serious, life-threatening, and sometimes fatal cases have been reported in the postmarketing setting in multiple sclerosis patients receiving natalizumab.

Laboratory confirmation in those cases was based on positive PCR for viral DNA in the cerebrospinal fluid. The duration of treatment with natalizumab prior to onset ranged from a few months to several years. Monitor patients receiving TYRUKO for signs and symptoms of meningitis and encephalitis. If herpes encephalitis or meningitis occurs, TYRUKO should be discontinued, and appropriate treatment for herpes encephalitis/meningitis should be administered.

Acute retinal necrosis (ARN) is a fulminant viral infection of the retina caused by the family of herpes viruses (eg. varicella zoster). ARN has been observed in patients being administered natalizumab and can be potentially blinding. Patients presenting with eye symptoms such as decreased visual acuity, redness and painful eye should be referred for retinal screening for ARN. Following clinical diagnosis of ARN, discontinuation of TYRUKO should be considered in these patients (see Adverse Reactions, Post Market Adverse Drug Reactions, Herpes).

Other Opportunistic Infections:

Physicians should be aware of the possibility that other opportunistic infections may occur during TYRUKO therapy (including patients without co-morbidities or concurrent therapy) and should include them in the differential diagnosis of infections that occur in natalizumab treated patients. Serious, life-threatening and sometimes fatal cases have been reported. If an opportunistic infection is suspected, dosing with TYRUKO is to be suspended until such infections can be excluded through further evaluations. If a patient receiving TYRUKO is diagnosed with an opportunistic infection, TYRUKO should only be restarted after the infection has been treated and if the benefit and risks of resuming TYRUKO therapy have been deemed to be acceptable by the treating physician.

Stopping TYRUKO Therapy:

If a decision is made to discontinue treatment with natalizumab, the physician needs to be aware that natalizumab remains in the blood, and has pharmacodynamic effects (e.g increased lymphocyte counts) for approximately 12 weeks following the last dose. Starting other therapies during this interval will result in a concomitant exposure to natalizumab. For medicinal products such as interferon and glatiramer acetate, concomitant exposure of this duration was not associated with safety risks in clinical trials. No data are available in MS patients regarding concomitant exposure with immunosuppressant medication. Use of these

medicinal products soon after the discontinuation of natalizumab may lead to an additive immunosuppressive effect. This should be carefully considered on a case-by-case basis, and a wash-out period of natalizumab might be appropriate. Short courses of steroids used to treat relapses were not associated with increased infections in clinical trials.

Reproductive Health: Female and Male Potential

It is not known whether natalizumab can affect reproductive potential (See 7.1.1 Pregnant Women)

Fertility

See 16 Non-Clinical Toxicology.

7.1 Special Populations

7.1.1 Pregnant Women

There are no adequate and well-controlled studies of natalizumab therapy in pregnant women. In premarketing clinical trials, the extent of exposure is very limited.

Thrombocytopenia and anaemia in infants born to women exposed to natalizumab during pregnancy were reported in the post-marketing setting. Monitoring for platelet counts, haemoglobin and hematocrit is recommended in neonates born to women exposed to natalizumab during pregnancy.

Because animal reproduction studies are not always predictive of human response, this drug should only be used during pregnancy if clearly needed. If a woman becomes pregnant while taking TYRUKO, discontinuation of TYRUKO should be considered.

In reproductive studies in monkeys and guinea pigs, there was no evidence of teratogenic effects or effects on survival or growth of offspring at doses up to 30 mg/kg (7 times the human clinical dose based on body weight comparison). In one of five studies that exposed monkeys or guinea pigs during pregnancy, the number of abortions in treated (30 mg/kg) monkeys was 33% vs. 17% in controls. No effects on abortion rates were noted in any other study. A study in pregnant cynomolgus monkeys treated at 2.3-fold the clinical dose demonstrated natalizumab-related changes in the fetus. These changes included mild anemia, reduced platelet count, increased spleen weights, and reduced liver and thymus weights associated with increased splenic extramedullary hematopoiesis, thymic atrophy and decreased hepatic hematopoiesis. In offspring born to mothers treated with natalizumab at 7-fold the clinical dose, platelet counts were also reduced. This effect was reversed upon clearance of natalizumab. There was no evidence of anemia in these offspring.

7.1.2 Breast-feeding

Natalizumab has been detected in human milk. Because of this and the potential for serious adverse reactions in nursing infants from natalizumab, a decision should be made whether to discontinue nursing or to discontinue the drug, taking into account the importance of the drug to the mother.

7.1.3 Pediatrics

Pediatrics (<18 years): Safety and effectiveness of natalizumab in pediatric MS patients have

not been established. Therefore, Health Canada has not authorized an indication for pediatric use (see 1.1 Pediatrics).

7.1.4 Geriatrics

Geriatrics (>65 years): Clinical studies of natalizumab did not include sufficient numbers of patients to determine whether they respond differently than younger patients.

8 Adverse Reactions

8.1 Adverse Reaction Overview

Serious adverse drug reactions most frequently reported during treatment with natalizumab in clinical trials were infections (3.2% vs. 2.6% placebo, including urinary tract infection [0.8% vs. 0.3%] and pneumonia [0.6% vs. 0%]); acute hypersensitivity reactions (1.1% vs. 0.3%, including anaphylaxis/anaphylactoid reaction [0.8% vs. 0%]); depression (1.0% vs. 1.0%, including suicidal ideation [0.6% vs. 0.3%]); and cholelithiasis (1.0% vs. 0.3%). See 7 Warnings and Precautions, Immune, Hypersensitivity Reactions.

The most frequently reported adverse events leading to discontinuation of natalizumab therapy were urticaria (1%) and other hypersensitivity reactions (1%). See 7 Warnings and Precautions, Immune, Hypersensitivity Reactions.

In clinical trials, cases of PML have been reported. PML can cause severe disability or death. See 7 Warnings and Precautions, Infections, Progressive Multifocal Leukoencephalopathy.

8.2 Clinical Trial Adverse Reactions

Clinical trials are conducted under very specific conditions. The adverse reaction rates observed in the clinical trials therefore, may not reflect the rates observed in practice and should not be compared to the rates in the clinical trials of another drug. Adverse reaction information from clinical trials may be useful in identifying and approximating rates of adverse drug reactions in real-world use.

In placebo-controlled trials in 1617 patients with multiple sclerosis treated with natalizumab, the incidence of common events was balanced between the natalizumab-treated patients and those who received placebo. Adverse events leading to discontinuation of therapy occurred in 5.8% of patients receiving natalizumab and in 4.8% of patients receiving placebo. Events are listed in Table 3 by body system and frequency of occurrence in the natalizumab group.

Table 3 All Adverse Events in Placebo-Controlled Studies of MS Occurring with Incidence ≥1.0% in Natalizumab Group and > 0.5% in Natalizumab Group than Placebo Group

	Natalizumab n = 1617 (%)	Placebo n = 1135 (%)
Blood and lymphatic system disorders		
Anemia	30 (1.9%)	14 (1.2%)

	Natalizumab n = 1617 (%)	Placebo n = 1135 (%)
Cardiovascular		
Tachycardia	23 (1.4%)	9 (0.8%)
Gastrointestinal		
Abdominal pain	75 (4.6%)	43 (3.8%)
General disorders and administration site conditions		
Fatigue	445 (27.5%)	305 (26.9%)
Oedema Peripheral	62 (3.8%)	25 (2.2%)
Chest Pain	58 (3.6%)	35 (3.1%)
Rigors	55 (3.4%)	12 (1.1%)
Weight decreased	27 (1.7%)	11 (1.0%)
Injury, poisoning, procedural complications		
Limb injury	38 (2.4%)	20 (1.8%)
Thermal burn	29 (1.8%)	12 (1.1%)
Immune system disorders		
Seasonal allergy	58 (3.6%)	35 (3.1%)
Infections and infestations		
Influenza	225 (13.9%)	146 (12.9%)
Sinusitis	184 (11.4%)	122 (10.7%)
Upper respiratory tract infection viral	134 (8.3%)	88 (7.8%)
Pharyngitis	125 (7.7%)	59 (5.2%)
Gastroenteritis	56 (3.5%)	21 (1.9%)
Tonsillitis	51 (3.2%)	23 (2.0%)
Bladder infection	38 (2.4%)	16 (1.4%)
Herpes zoster	33 (2.0%)	16 (1.4%)
Respiratory tract infection	30 (1.9%)	15 (1.3%)
Gingival infection	18 (1.1%)	6 (0.5%)
Musculoskeletal and connective tissue disorders		
Muscle Cramp	82 (5.1%)	42 (3.7%)
Joint swelling	32 (2.0%)	13 (1.1%)
Nervous system disorders		
Headache	634 (39.2%)	436 (38.4%)
Dysesthesia	42 (2.6%)	23 (2.0%)
Sinus headache	38 (2.4%)	19 (1.7%)
Psychiatric disorders		
Depressed mood	37 (2.3%)	16 (1.4%)
Reproductive system and breast disorders		
Menstruation irregular	37 (2.3%)	12 (1.1%)

	Natalizumab n = 1617 (%)	Placebo n = 1135 (%)
Respiratory, thoracic and mediastinal disorders		
Cough	130 (8.0%)	81 (7.1%)
Sinus congestion	51 (3.2%)	22 (1.9%)
Epistaxis	28 (1.7%)	13 (1.1%)
Vascular disorders		
Hematoma	17 (1.1%)	6 (0.5%)

Additional Information:

Hypersensitivity:

In post-marketing experience, there have been reports of hypersensitivity reactions with hypotension, hypertension, chest pain, chest discomfort, dyspnoea, and angioedema, in addition to more usual symptoms such as rash and urticaria.

The incidence of hypersensitivity reactions was based on the investigator assessment that the event was urticaria or an allergic reaction, which may have included terms such as urticaria, itch, flushing, hypersensitivity or anaphylactoid reaction. In controlled clinical trials in MS patients, hypersensitivity reactions occurred in up to 4% of patients. Serious systemic hypersensitivity reactions (e.g. anaphylactic/anaphylactoid) occurred in <1% (study 1801: 5/627) of MS patients. Hypersensitivity reactions usually occurred within two hours of the start of the infusion.

Immunogenicity:

Persistent anti-natalizumab antibodies (detected on two occasions at least 6 weeks apart) were associated with decreased efficacy of natalizumab and an increased incidence of hypersensitivity reactions. The majority of patients who became persistently antibody-positive had developed antibodies by 12 weeks.

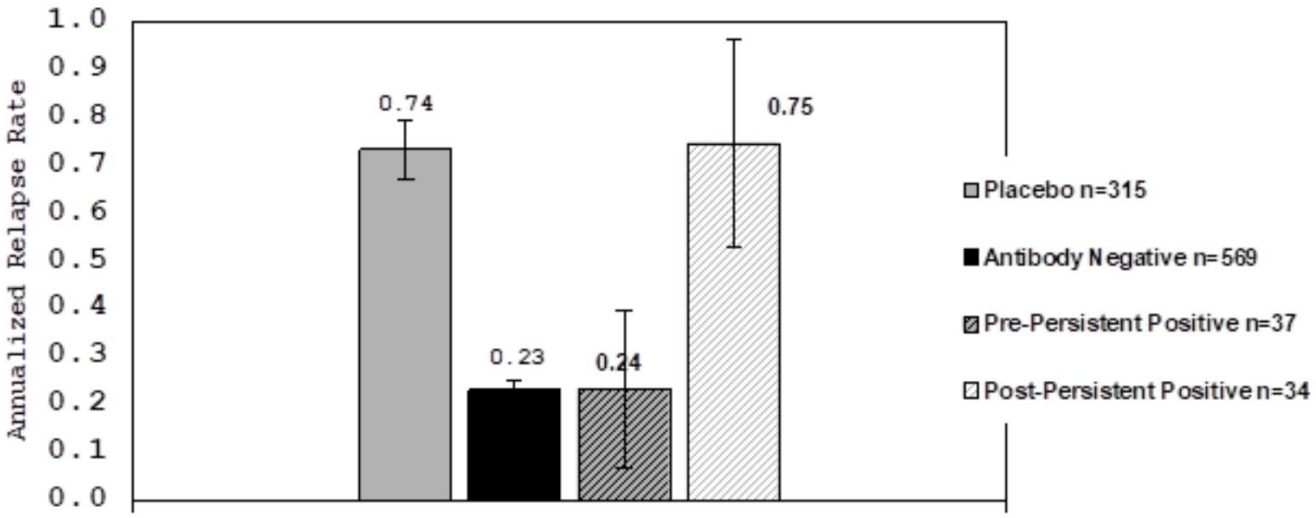
In controlled clinical trials in MS patients, persistent anti-natalizumab antibodies developed in approximately 6% of patients. Antibodies were detected on only one occasion in 4% of patients. Additional infusion-related reactions associated with persistent antibodies included rigors, nausea, vomiting and flushing. Approximately 90% of patients who became persistently antibody-positive in 2-year clinical trials had developed antibodies by 12 weeks.

If, after 3 months of natalizumab treatment, the presence of persistent antibodies is suspected, or in patients who have received an initial short exposure to natalizumab and extended periods without treatment, antibody testing should be performed. Antibodies may be detected and confirmed with sequential serum antibody tests. Antibodies detected early in the treatment course (e.g. within 6 months) may be transient and disappear with continued dosing. Repeat testing between 6 weeks and 3 months after the initial positive result is recommended in patients in whom antibodies are detected to confirm that antibodies are persistent. In the presence of persistent antibodies, discontinuation of treatment with TYRUKO should be considered (see Figure 2).

Patients who receive natalizumab for a short exposure (1-2 doses) followed by an extended period without treatment are at higher risk of developing anti-natalizumab antibodies and/or hypersensitivity reactions on re-exposure. Given that patients with persistent antibodies to natalizumab experienced reduced efficacy and that hypersensitivity reactions were more common in such patients, consideration should be given to testing for the presence of persistent antibodies prior to redosing following a prolonged dose interruption.

Information regarding the availability and location of testing laboratories may be obtained by contacting Sandoz PLUS at (1-888-449-7673).

Figure 2: Subject Relapse Rate Prior to and After Antibody Detection – Persistent Positives – Study 1801.



Infections:

In controlled clinical trials in MS patients, the rate of infection was approximately 1.5 per patient year in both natalizumab and placebo-treated patients. The nature of the infections was generally similar in natalizumab and placebo-treated patients. The majority of patients did not interrupt natalizumab therapy during infections, and recovery occurred with appropriate treatment.

In clinical trials, cases of PML have been reported (see Warnings and Precautions, Infections; Adverse Drug Reaction Overview).

In other clinical trials, cases of opportunistic infections have been reported, some of which were fatal. While a causal role for natalizumab cannot be excluded, it is reasonable to conclude that comorbidities and concomitant medications played an important role in these infections. Should a serious opportunistic infection develop, natalizumab therapy should be withheld until the infection has been successfully treated.

TYRUKO should only be restarted after the infection has been treated and if the benefit and risks of resuming TYRUKO therapy have been deemed to be acceptable by the treating physician (see Warnings and Precautions, Infections, Other Opportunistic Infections).

In clinical trials, herpes infections occurred slightly more frequently in natalizumab-treated patients than in placebo-treated patients. During post-marketing experience, there have been

rare reports of serious cases including encephalitis and meningitis; some cases have been life-threatening and sometimes fatal. Should a serious herpes infection occur, natalizumab therapy should be withheld until the infection has been treated.

Short courses of corticosteroids can be used in combination with natalizumab. In phase 3 MS clinical trials, concomitant treatment of relapses with a short course of corticosteroids was not associated with an increased rate of infection in patients treated with natalizumab as compared with those on placebo.

Infusion-Related Reactions:

An infusion-related reaction was defined in clinical trials as any adverse event occurring within 2 hours of the start of an infusion. These events occurred in 23.1% of MS patients treated with natalizumab (18.7% placebo). Events reported more commonly with natalizumab than with placebo included headache, dizziness, fatigue, urticaria, pruritus and rigors.

Malignancies:

No differences in incidence rates or the nature of malignancies between natalizumab and placebo- treated patients were observed over 2 years of treatment. However, observation over longer treatment periods is required before any effect of natalizumab on malignancies can be excluded. Should a malignancy develop, natalizumab therapy should be withheld at least until appropriate treatment has been initiated for the malignancy and the benefit and risks of resuming natalizumab therapy have been deemed to be acceptable by the treating physician.

8.3 Less Common Clinical Trial Adverse Reactions

The incidence of adverse drug reactions experienced by <1% of subjects in natalizumab group and at least 0.1% higher in natalizumab compared to placebo are listed below:

Blood and lymphatic system disorders: Anemia, thrombocytopenia, leukocytosis

Cardiac disorders: Tachycardia, angina pectoris

Ear and labyrinth disorders: Vertigo

Gastrointestinal disorders: Flatulence, upper abdominal pain, abdominal distention, epigastric discomfort

General disorders and administration site conditions: Feeling hot, peripheral edema, lethargy, feeling abnormal, infusion site erythema, pain, thirst, hyperpyrexia, infusion site pruritus

Immune system disorders: Hypersensitivity, anaphylactoid reaction, anaphylactic reaction

Infections and infestations: Progressive multifocal leukoencephalopathy, pharyngitis, sinusitis, herpes simplex, herpes zoster, rhinitis infective, bronchial infection, gastroenteritis, skin and subcutaneous tissue abscess, furuncle, pharyngitis streptococcal, bladder infection, breast abscess, dermatitis infected, herpes viral infection, oral infection, pharyngitis viral, tooth infection, urinary tract infection

Injury, poisoning and procedural complications: Overdose

Investigations: Aspartate aminotransferase increased, neutrophil count increased, heart rate

increased, neutrophil count decreased, white blood cell count increased, blood test abnormal

Musculoskeletal and connective tissue disorders: Myalgia, muscle cramp, muscle spasms, sensation of heaviness, joint stiffness, muscle tightness, muscle weakness

Neoplasms benign, malignant and unspecified (incl cysts and polyps): Cyst

Nervous system disorders: Tremor, paresthesia oral, sensory disturbance, paresis, psychomotor hyperactivity, syncope

Psychiatric disorders: Depression, agitation

Reproductive system and breast disorders: Irregular menstruation

Respiratory, thoracic and mediastinal disorders: Cough, sinus congestion, wheezing, throat irritation

Skin and subcutaneous tissue disorders: Erythema, rash pruritic, acne, pruritus, urticaria, dry skin, onychorrhexis, skin irritation

Vascular disorders: Petechiae, poor venous access, thrombophlebitis, vasodilatation

8.5 Post-Market Adverse Reactions

The following adverse reactions have been identified during post-marketing use of natalizumab. Because these reactions are reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate their frequency or to establish a causal relationship to drug exposure.

Hypersensitivity: There have been reports of hypersensitivity reactions which have been associated with one or more of the following: hypotension, hypertension, chest pain, chest discomfort, and dyspnea.

Hepatic: There have been rare reports of clinically significant liver injury, including markedly elevated serum hepatic enzymes and elevated total bilirubin, as early as 6 days after the first dose. In some patients, liver injury recurred upon rechallenge. Some cases occurred in patients with pre-existing liver disease or in the presence of other drugs that have been associated with hepatic injury.

Infections:

Progressive Multifocal Leukoencephalopathy:

There have been rare reports of progressive multifocal leukoencephalopathy (PML) in patients with MS receiving natalizumab monotherapy, including cases with onset in the absence of clinical symptoms of PML. Some cases have been reported up to 6 months following discontinuation of natalizumab monotherapy (see Warnings and Precautions, Infections, Progressive Multifocal Leukoencephalopathy).

Patients who have all three risk factors for PML (i.e. anti-JCV antibody positive AND have received more than 2 years of natalizumab therapy AND have received prior immunosuppressant therapy) are at a higher risk of PML.

Anti-JCV antibody testing may provide supportive information for PML risk stratification prior to or during treatment with TYRUKO.

Cases of JCV GCN have also been reported during postmarketing use of natalizumab.

Herpes: There have been rare reports of serious cases of herpes infections.

In post marketing experience, acute retinal necrosis (ARN) has been observed at a higher incidence in patients receiving natalizumab. Some cases have occurred in patients with central nervous system (CNS) herpes infections (eg. herpes meningitis and encephalitis). Serious cases of ARN, either affecting one or both eyes, led to blindness in some patients. The treatment reported in these cases included anti-viral therapy and in some cases, surgery.

Hematologic: In post-marketing experience, there have been reports of eosinophilia (eosinophil count > 1,500/mm^3) without clinical findings. In cases where natalizumab therapy was discontinued the elevated eosinophil levels resolved. Rare serious cases of anemia and hemolytic anemia have been reported in patients treated with natalizumab in post-marketing observational studies.

Cases of thrombocytopenia, some of which were serious, in infants born to women exposed to natalizumab during pregnancy were reported in the postmarketing setting. Cases of anemia were reported in infants born to women exposed to natalizumab. Therefore, monitoring for hematological abnormalities is recommended in neonates who were exposed to natalizumab *in utero* (see 7.1.1 Pregnant Women).

In post-marketing experience, there have been reports of uncommon frequency of thrombocytopenia and immune thrombocytopenic purpura (ITP), some of which were serious (see 7 Warnings and Precautions, Hematologic).

9 Drug Interactions

9.2 Drug Interactions Overview

If a decision is made to discontinue treatment with TYRUKO, the physician needs to be aware that natalizumab has pharmacodynamic effects (e.g. increased lymphocyte counts) for approximately 12 weeks following the last dose. For drugs such as interferon and glatiramer acetate, concomitant exposure of this duration was not associated with safety risks in clinical trials. This should be carefully considered on a case-by-case basis and a washout period of TYRUKO might be appropriate.

Should TYRUKO therapy be administered after treatment with another immunosuppressive drug, physicians should consider the half-life of the drug and the potential for persistent immunosuppressive effects of these products when considering if a washout period is needed and, if so, its duration.

TYRUKO should not be diluted with anything other than 0.9% Sodium Chloride Injection, USP.

9.3 Drug-Behavioural Interactions

Drug-Behavioural Interactions have not been established.

9.4 Drug-Drug Interactions

Interactions with other drugs have not been established.

9.5 Drug-Food Interactions

Interactions with food have not been established.

9.6 Drug-Herb Interactions

Interactions with herbal products have not been established.

9.7 Drug-Laboratory Test Interactions

In clinical trials, natalizumab was observed to induce increases in circulating lymphocytes, monocytes, eosinophils and nucleated red blood cells. Observed increases persisted during natalizumab exposure, but were reversible, returning to baseline levels usually within 16 weeks after the last dose. Elevations of neutrophils were not observed.

Natalizumab may induce mild decreases in hemoglobin levels (mean decrease of 6.0 g/L). Hemoglobin levels returned to pretreatment values, usually within 16 weeks of last dose of natalizumab and the changes were not associated with clinical symptoms.

10 Clinical Pharmacology

10.1 Mechanism of Action

Natalizumab is a selective adhesion molecule (SAM) inhibitor and binds to the α4-subunit of human integrin, which is highly expressed on the surface of all leukocytes, with the exception of neutrophils.

Specifically, natalizumab binds to the $\alpha4\beta1$ integrin blocking the interaction with its cognate receptor, vascular cell adhesion molecule-1 (VCAM-1), and additional ligands such as osteopontin, and an alternatively spliced domain of fibronectin, connecting segment-1 (CS-1). Natalizumab blocks the interaction of $\alpha4\beta7$ integrin with the mucosal addressin cell adhesion molecule-1 (MadCAM-1). Disruption of these molecular interactions prevents transmigration of mononuclear leukocytes across the endothelium into inflamed parenchymal tissue. A further mechanism of action of natalizumab may be to suppress ongoing inflammatory reactions in diseased tissues by inhibiting the interaction of α4-expressing leukocytes with their ligands in the extracellular matrix and on parenchymal cells. As such, natalizumab may act to suppress inflammatory activity present at the disease site, and inhibit further recruitment of immune cells into inflamed tissues.

In multiple sclerosis (MS), lesions are believed to occur when activated inflammatory cells, including T-lymphocytes, cross the blood-brain barrier (BBB). Leukocyte migration across the BBB involves interaction between adhesion molecules on inflammatory cells and endothelial cells of the vessel wall. The interaction between $\alpha4\beta1$ and its targets is an important component of pathological inflammation in the brain, and disruption of these interactions leads to reduced inflammation. Under normal conditions, VCAM-1 is not expressed in the brain parenchyma.

However, in the presence of pro-inflammatory cytokines, VCAM-1 is upregulated on endothelial cells, and possibly on glial cells near the sites of inflammation. In the setting of central nervous system (CNS) inflammation in MS, it is the interaction of $\alpha4\beta1$ with VCAM-1, CS-1 and osteopontin that mediates the firm adhesion and transmigration of leukocytes into the brain parenchyma, and may perpetuate the inflammatory cascade in CNS tissue. Blockade of the molecular

interactions of $\alpha 4\beta 1$ with its targets reduces inflammatory activity present in the brain in MS and inhibits further recruitment of immune cells into inflamed tissue, thus reducing the formation or enlargement of MS lesions.

10.2 Pharmacodynamics

Treatment with natalizumab led to an increase in circulating white blood cells and total lymphocytes that was maintained throughout the treatment period. This is due to the ability of natalizumab to inhibit adhesion of leukocytes to endothelial cells and diminish transmigration of these cells from the vascular space into inflamed tissues. These increases were not clinically significant and once treatment was discontinued, counts returned to baseline levels. Consistent with the mechanism of action of natalizumab and the lack of $\alpha 4$ on the surface of this cell type, there was no change in the number of circulating neutrophils.

In study 101MS101, $\alpha 4$-integrin saturation levels were generally consistent with those observed in the phase 3 studies (C-1801 and C-1802). In these studies which demonstrated efficacy, $\alpha 4$-integrin saturation approximating 70% was observed, however there was a high degree of assay variability.

10.3 Pharmacokinetics

Pharmacokinetic values determined after a single 300 mg dose of natalizumab in healthy subjects are provided in Table 4. Similar values observed in MS patients after a single dose and after 6 months of dosing as monotherapy are given in Table 5. Some accumulation occurs over the 6- month dosing period.

Table 4: Pharmacokinetic Parameters, Single-Dose 300 mg Natalizumab as Intravenous Infusion of 60 minutes

Median Values of Parameter	Study 1805	Study 1806
AUCτ (μg/mL *hr)	19900	21500
Cmax (μg/mL)	110	94
Tmax (hrs)	2.98	3.00
t ½ (hr)	224	249
Vdis (mL/kg)	66.6	67.4
CL (mL/hr/kg)	0.212	0.179

Table 5: Summary of Pharmacokinetic Parameters Following 60-Minute 300 mg Natalizumab Infusions Given Monthly in MS Patients (Mean +/- s.d.)

Dose Number	Study	C_{max} (μg/mL)	Minimum (Trough) Conc. (μg/mL)	$AUC_{(last)}$ (μg×hr/mL)	Vd (mL/kg)	CL (mL/hr/kg)	$t_{½}$ (hr)

1	C-1801	84.8 ± 22.3	none	17884 ± 9165	77 ± 36	0.23 ± 0.09	249 ± 105
6	C-1801	94.7 ± 34.2	21.3 ± 15.3*	19609 ± 5701	81 ± 43	0.22 ± 0.06	265 ± 98

*Representative of concentration at the end of 6-months dosing (24-week measurement)

Clinical studies, 101MS101 and 101MS102, demonstrated that the pharmacokinetic, pharmacodynamic, immunogenicity and safety profiles of drug product produced from a high yield drug substance manufacturing process are comparable to the drug product produced from the original drug substance manufacturing process.

Special Populations and Conditions

- **Pediatrics:** The pharmacokinetics of natalizumab in pediatric MS patients have not been established.
- **Geriatrics:** The pharmacokinetics of natalizumab in MS patients over 65 years of age have not been established.
- **Sex:** Results of a population pharmacokinetic demonstrated that gender did not influence natalizumab pharmacokinetics.
- **Ethnic Origin:** The effects of race on the pharmacokinetics of natalizumab have not been studied.
- **Hepatic Insufficiency:** The pharmacokinetics of natalizumab in patients with hepatic insufficiency have not been studied.
- **Renal Insufficiency:** The pharmacokinetics of natalizumab in patients with renal insufficiency have not been studied.

Duration of Effect:

Natalizumab has pharmacodynamic effects (e.g. increased lymphocyte counts) for approximately 12 weeks following the last dose.

11 Storage, Stability And Disposal

TYRUKO (natalizumab for injection) single-use vials must be stored in a refrigerator between 2°C to 8°C. Do not use beyond the expiration date on the carton and vial label. Do not shake or freeze. Protect from light.

TYRUKO does not contain preservatives, therefore reconstitution and dilution of the product should be performed under aseptic conditions.

After dilution in 0.9% sodium chloride solution, if not used immediately, store the TYRUKO solution for infusion at 2°C to 8°C. The administration of TYRUKO solution for infusion must be completed within 24 hours of dilution.

Any unused product remaining in the vials and/or waste material should be disposed of in accordance with local requirements.

12 Special Handling Instructions

TYRUKO (natalizumab for injection) is for single use only. One vial of TYRUKO should be diluted only with 0.9% Sodium Chloride Injection, USP before use.

Part II: Scientific Information

13 Pharmaceutical Information

Drug Substance

Proper name: natalizumab

Chemical name: Recombinant, humanized, anti-α4–integrin monoclonal antibody
Molecular formula and molecular mass: Approximately 149 kilodaltons.

Physicochemical properties: Natalizumab is produced by recombinant DNA technology in a Chinese Hamster Ovary mammalian cell line. Natalizumab is of the IgG_4 subclass and consists of two heavy and two light chains connected by four inter-chain disulfide bonds.

Product Characteristics:

TYRUKO (natalizumab for injection) is a recombinant humanized $IgG_{4κ}$ monoclonal antibody selective for α4- integrin. Natalizumab is produced in Chinese Hamster Ovary mammalian cells. The molecular weight of natalizumab is 149 kilodaltons. TYRUKO is supplied as a sterile, colourless, clear to slightly opalescent concentrate for solution for intravenous (IV) infusion.

14 Clinical Trials

14.1 Clinical Trials by Indication

Multiple Sclerosis (MS):

Table 6 - Summary of patient demographics for clinical trials in MS

Study #	Study design	Dosage, route of administration and duration	Study subjects (n)	Median age (Range)	Sex
Pivotal Studies					
1801 AFFIRM	Randomized, double-blind, placebo-controlled study. MS patients who had experienced at least one clinical relapse during the prior year. EDSS score between 0.0 and 5.0	300 mg IV or placebo every 4 weeks for up to 30 infusions	942 (627 natalizumab and 315 placebo)	37 (18-50)	F: 660 (70%) M: 282 (30%)

1802 SENTINEL	Randomized, double-blind, placebo-controlled study. MS patients who had experienced at least one clinical relapse during the prior year on therapy with Avonex 30 µg intramuscularly (IM) once weekly. EDSS score between 0.0 and 5.0	300 mg IV or placebo in combination with Avonex 30 µg IM once weekly every 4 weeks for up to 30 infusions	1171 (589 natalizumab and 582 in combination with Avonex)	39 (18-55)	F: 862 (74%) M: 309 (26%)
Supportive Studies					
MS-231	Randomized, double-blind, placebo-controlled multicentre study in patients with relapsing remitting MS or secondary progressive MS	3 mg/kg, 6 mg/kg or placebo every 28 days for 6 infusions	213 (71 placebo, 68 natalizumab 3 mg/kg, 74 natalizumab 6 mg/kg)	44 (22-66)	F: 152 (71.4%) M: 61 (28.6%)

Study Results:

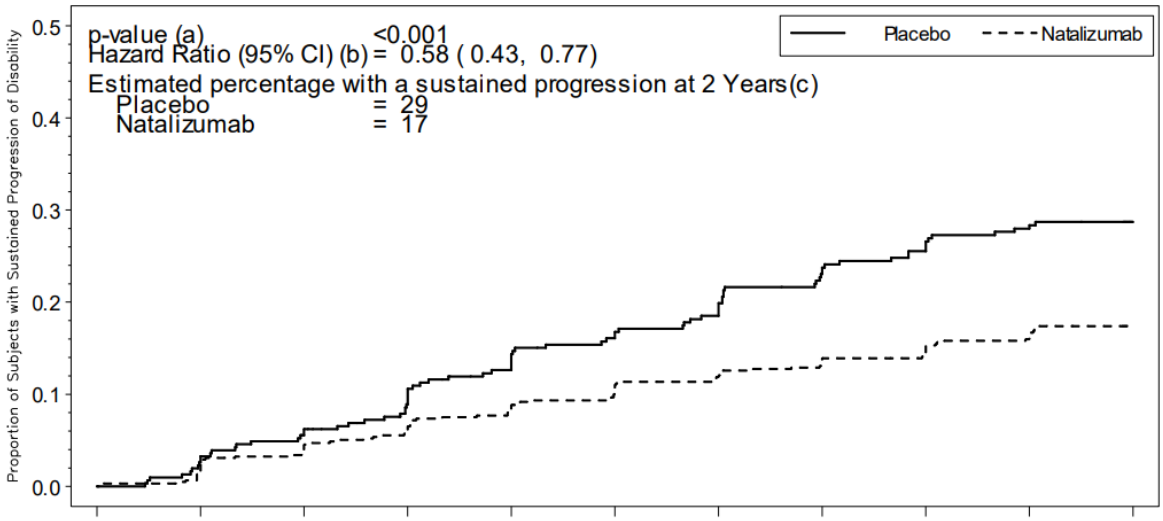
Table 7 - Results of study 1801 (AFFIRM) in MS

	natalizumab* n=627	**Placebo*** n=315	***p*value**
<u>Clinical Endpoints</u>			
Percentage with sustained progression of disability (increase in EDSS sustained for 12 weeks)	17%	29%	
Hazard ratio	0.58, (95% CI 0.43, 0.77)		$p<0.001$
Risk reduction	42%		

Percentage with sustained progression of disability (increase in EDSS sustained for 24 weeks) Hazard ratio Risk reduction	11% 0.46 (95% CI 0.33, 0.64) 54%	23%	*p*<0.001
Annualized relapse rate (Percent reduction compared with placebo)	0.24 (68%)	0.73	*p*<0.001
Percentage of patients relapse-free after 2 years	67%	41%	$p<0.001$
MRI Endpoints			
Median percentage change in volume of T2-hyperintense lesions	-9.4%	8.8%	*p*<0.001
Number of new or newly-enlarging T2-hyperintense lesions Mean Percent reduction compared with placebo	1.9 83%	11.0	*p*<0.001
Median Percentage of patients with: 0 lesions 1 lesion 2 lesions 3 or more lesions	0.0 57% 17% 8% 18%	5.0 15% 10% 8% 68%	*p*<0.001
Number of Gd-enhancing Lesions Mean Percent reduction compared with placebo	0.1 92%	1.2	*p*<0.001
Median Percentage of patients with: 0 lesions 1 lesion 2 or more lesions	0.0 97% 2% 1%	0.0 72% 12% 16%	*p*<0.001

*All analyses were intent-to-treat.

Figure 3: Time to Sustained Progression of Disability as Measured by Increase in EDSS in AFFIRM (1801)



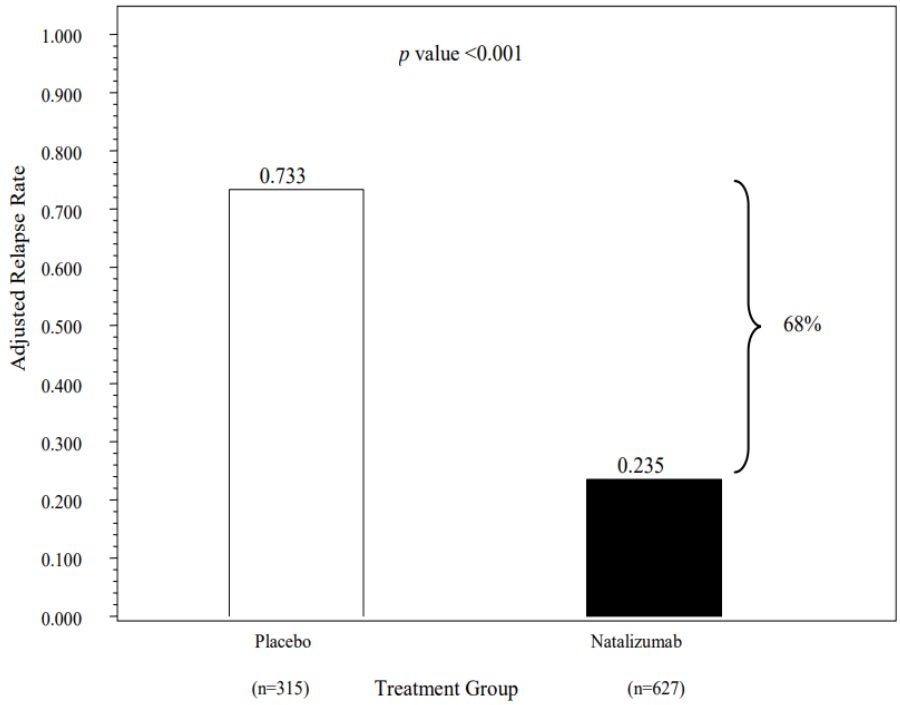
Number of Patients at Risk

Placebo	315	296	283	264	248	240	229	216	208	200
Natalizumab	627	601	582	567	546	525	517	503	490	478

NOTE: Sustained progression of disability is defined as at least 1.0 point increase on the EDSS from a baseline EDSS >=1.0 sustained for 12 weeks or at least a 1.5 point increase on the EDSS from a baseline EDSS of 0 sustained for 12 weeks.

(a) Log Rank p-value.
(b) Hazard ratio (natalizumab/placebo) estimated from a Cox proportional hazards model adjusting for baseline EDSS and age (<40 versus >=40)
(c) Kaplan Meier estimate of the percentage of subjects expected to have sustained progression within 2 Years

Figure 4: Annualized Relapse Rates in AFFIRM (1801)



Comparing studies 1801 and 1802, at baseline, those patients in 1802 had, on average, longer disease duration and higher EDSS scores than patients in study 1801.

The efficacy of natalizumab alone was not compared with the efficacy of natalizumab plus Avonex.

Other Clinical Endpoints:

Brain Atrophy:

Brain parenchymal fraction (BPF) is a marker of the destructive pathologic process ongoing in relapsing MS patients, with changes measuring a loss of brain tissue.

In the monotherapy study, 1801, the BPF interval changes at 1 year and 2 years suggests a pattern that differs between the placebo and treated groups. Although both groups exhibited a decrease in BPF from baseline to year 1, there was a relatively greater reduction in the natalizumab group compared to the placebo group (natalizumab vs. placebo: 0.56% vs. 0.40%, p=0.002). Between 1 and 2 years, the placebo group again demonstrated an equivalent reduction in brain volume (0.43%) comparable to year 1 while the reduction observed in the natalizumab-treated group decreased to 0.24% indicating a significant slowing of atrophy compared to the placebo group (p= 0.004).

From baseline to 1 year the placebo- and natalizumab-treated groups had a decreased BPF during the first year with a relatively greater reduction in the natalizumab treated groups. The disproportionate reduction in the natalizumab-treated groups during the first year is likely due to an acute reduction in inflammation and edema. The lower rate of atrophy seen in the second year of each study likely reflects the significant anti-inflammatory treatment effects of natalizumab on atrophy progression.

Hospitalization:

The number of hospitalizations was recorded in phase 3 studies. Table 8 summarizes the number of hospitalizations whose primary reason was categorized as MS relapse, MS-related complication, elective surgery or "other". The number of hospitalizations for MS-related or MS relapses, reported separately, followed a similar pattern to all hospitalizations.

Table 8: Rate of Hospitalization

	AFFIRM (1801)		
	Placebo	**Natalizumab**	
Number of subjects with a hospitalization (%)	77 (24%)	114 (18%)	
Total number of hospitalizations	125	156	
Annualized hospitalization rate (95% CI)	0.183 (0.142, 0.236)	0.112 (0.089, 0.140)	Rate ratio 0.612 (0.436, 0.858) p=0.005
Number of subjects with MS-related hospitalizations (%)	41 (13%)	37 (6%)	
Total number of MS-related hospitalizations	66	48	
Annualized MS-related hospitalization rate (95% CI)	0.097 (0.070, 0.133)	0.034 (0.024, 0.050)	Rate ratio 0.356 (0.218, 0.582) p<0.001

Quality of Life (QOL):

The Multiple Sclerosis Quality of Life Index (MSQLI) consists of several scales to measure important symptoms experienced by MS patients as part of their disease. Patients treated with natalizumab alone showed on average an increase in both the physical and mental component scores of the SF-36 indicating improvement, whereas those who received placebo showed worsening on average ($p \leq 0.01$). No statistically significant differences were observed between treatment groups on the non-SF-36 components. Indeed, there were minimal changes in either group on many of these measures.

Patients were asked how they were feeling during the study and to mark this on a 10 cm Visual Analogue Scale (VAS), with 0 as "poor" and 100 as "excellent". On average, patients treated with natalizumab had a significant effect on sense of well-being vs. patients treated with placebo using the VAS (p=0.007).

Subgroup Analyses of Efficacy:

Results for disability progression and clinical relapses have been tabulated by baseline EDSS (≤3.5, >3.5), the number of T2 lesions at baseline (<9, ≥9), presence or absence of Gd-lesions and age at baseline (<40, ≥40 years).

Natalizumab consistently delayed disability progression and reduced relapse rates in most clinical subpopulations, even those with the highest degree of disease activity. Although the subgroup of patients with fewer than 9 T2 lesions at baseline showed a treatment effect on

the secondary MRI outcome measures, they did not show a similar effect on the primary clinical outcome measures; this may be due to the small numbers in this group. To the contrary, a significant effect was seen in those subjects with 9 or more T2 lesions at baseline. Patients with MS defined by McDonald diagnostic criteria 1 through 4 demonstrated consistent clinical responses to natalizumab, although the subgroup defined by McDonald diagnostic criteria 1 or 2 appeared to have a greater response than criteria 3 or 4. These data indicate that the degree of MS disease activity should be a consideration when starting natalizumab treatment.

There were twice as many women as men in study 1801. Treatment with natalizumab was equally effective for both women and men across all 2-year endpoints.

Using the Multiple Sclerosis Functional Composite Score (Timed 25-Foot Walk [T25FW], 9 Hole

Peg Test [9HPT], and the Paced Auditory Serial Addition Test 3 [PASAT3]), in study 1801, the group treated with natalizumab showed statistically significant improvement in the overall score (p<0.0001), and on each parameter (T25FW: p<0.0001, 9HPT: p<0.0001, PASAT3: p=0.005) compared to the placebo-treated group.

In study 1801, there was a statistically significant relative reduction (69% p<0.001) in the rate of relapses treated with steroids (43% placebo-treated group vs. 13% of the natalizumab-treated group experienced relapses).

Phase 2 Clinical Study

In a randomized, double-blind phase 2 trial (MS-231), a total of 213 patients with relapsing-remitting (RRMS, n=144) and relapsing secondary progressive (SPMS, n=69) multiple sclerosis received either placebo (n=71) or natalizumab 3 mg/kg/body weight (n=68) or 6 mg/kg/body weight (n=74) every 28 days for 6 months. The primary endpoint was the number of new brain lesions on monthly gadolinium-enhanced magnetic resonance imaging during the 6-month treatment period. Clinical outcomes included relapses and self-reported well-being. The results indicated that both groups treated with natalizumab had fewer enhancing lesions compared to the placebo- treated group. This difference was statistically significant (p<0.001). A similar result was noted for the number of patients with a relapse (p= 0.02), or with an objective relapse (p=0.004), or requiring corticosteroid treatment (p <0.001). Overall patients treated with natalizumab had an improved sense of well-being using the Visual-Analog Score. During this study natalizumab was well tolerated with a safety profile similar to that of placebo. Headache was the most common reported adverse event. The clinical significance of a trend toward an increased rate of infection in the natalizumab-treated patients was unclear.

14.3 Immunogenicity

In the 2-year studies with monthly administration, continued measurements showed that the incidence of anti- natalizumab antibodies at a single timepoint was 10%, 6% being persistent anti-natalizumab antibodies (detected on two occasions at least 6 weeks apart) and 4% transient.

Persistent anti-natalizumab antibodies were associated with a decreased efficacy of natalizumab and an increased incidence of infusion-related reactions. The majority of patients who became persistently antibody-positive had developed antibodies by 12 weeks. Persistent antibody-positive subjects had trough natalizumab concentrations that were consistently below limit of quantitation. This level of natalizumab exposure was correlated with consistent reduction in α-4 integrin saturation on leukocytes in these persistently antibody-positive subjects. Patients who experienced transient anti- natalizumab antibodies had a temporary reduction in natalizumab concentrations, but did *not* experience a long-term reduction in drug concentrations or activity.

16 Non-Clinical Toxicology – Reference Biologic Drug

General Toxicology: Natalizumab is a human IgG_4 molecule and is therefore a foreign protein and immunogenic in the various species used in the toxicology program. The rate and incidence of detectable antibody formation varies with species, dose level and dosing regimen. The development of antibodies can affect the detection of natalizumab levels and can also reduce exposure to the drug through accelerated clearance. Therefore toxicokinetic sampling was performed in the majority of the chronic toxicology studies to allow for the monitoring of exposure over time.

Natalizumab treatment has been generally well tolerated in both adult and juvenile animals at cumulative exposures in the high-dose group (60 mg/kg) of up to 94-fold (mean 36, range 0.4-94) and 78-fold (mean 53, range 36-78), respectively, over anticipated human exposures (based on 6 months of dosing and a human dose of 300 mg).

In all species tested, increases in white blood cell count were associated with serum natalizumab levels above approximately 1-5μg/mL and returned to normal when natalizumab levels fell below these values. Increases were primarily the result of increases in lymphocyte counts, though increases in monocytes, eosinophils and basophils were also seen. These increases are the expected pharmacologic effect of the binding of natalizumab to α4-integrins, as α4 integrins are involved in the adhesion of leukocytes to endothelium and the subsequent trafficking of the leukocytes across the endothelium and into tissues.

Findings (some seen inconsistently across the studies) other than increased WBC counts included: increases in reticulocytes and/or nucleated red blood cells, increased spleen weights, mild to moderate follicular hyperplasia in the lymph node and spleen, and minimal to mild focal leukocyte infiltrates in the liver. Hyperplastic lymphoid follicles retained normal anatomic relationships and boundaries and were considered characteristic of follicles exhibiting polyclonal lymphocyte expansion and/or accumulation. Findings have been reversible following the discontinuation of natalizumab treatment.

Overall, natalizumab-related treatment findings seen in the toxicology program were generally

related (e.g., hematology changes) or believed to be related (e.g., increased spleen weights) to the pharmacological activity of natalizumab. Treatment-related effects were not associated with overt toxicity in the affected animals, and were reversible following discontinuation of natalizumab treatment and clearance of natalizumab from circulation.

Genotoxicity: Natalizumab was not genotoxic in a human lymphocyte aberration assay, nor did it have any effect on the proliferation of α4-expressing human tumour cells lines in vitro. Growth and metastasis of two α4-expressing tumours (a leukemia and a melanoma) were not increased in the presence of natalizumab in nude and SCID mouse human tumour xenograft models.

Reproductive and Developmental Toxicology: Male fertility in guinea pigs was unaffected by natalizumab treatment at doses resulting in cumulative AUC exposures 38-fold times that in humans. Female fertility in guinea pigs demonstrated treatment-related effects (consisting of reduced fertility and reduced pup survival through post-natal day [PND] 14) at doses resulting in cumulative AUC exposures 40-fold times that in humans – effects were not observed at the next lowest dose level that resulted in cumulative AUC exposures 13-fold times that in humans. No teratogenic effects in guinea pigs or monkeys were seen as a result of natalizumab treatment. There was an apparent increased incidence of abortion in natalizumab-treated primates in only one of four studies that evaluated this effect, in guinea pigs and cynomolgus monkeys. Increases in nucleated red blood cells, changes in RBC parameters indicative of mild anemia, and some alterations in the distribution of immune cell populations in immune system organs were seen in the fetuses of guinea pigs and/or primates at high exposure levels (mean cumulative AUC 65-fold over human). Although no effects were noted at the lowest dose tested, exposure at this dose was only achieved through GD44. In offspring born to mothers treated with natalizumab at 7-fold the clinical dose, reduced platelet counts were observed and were reversed upon clearance of natalizumab. There was no evidence of anemia in these offspring. Offspring exposed in utero and via breast milk had no natalizumab-related changes in the lymphoid organs and had normal immune response to challenge with a T-cell dependent antigen.

Tissue cross-reactivity studies of natalizumab with normal adult human, cynomolgus and rhesus monkey and guinea pig tissues, and with fetal human and monkey tissues were consistent with staining patterns reported in the literature and did not result in the identification of any unexpected target tissues. Findings in the chronic and reproductive toxicology studies involved some of the lymphoid organs seen to stain in these studies, but did not reveal any findings associated with the less common, nonlymphoid staining patterns.

17 Supporting Product Monographs

1. PrTYSABRI®, concentrate for solution for intravenous infusion 300 mg/15 mL, submission control: 290559, Product Monograph, Biogen Canada Inc. (JAN 9, 2025).

Patient Medication Information

READ THIS FOR SAFE AND EFFECTIVE USE OF YOUR MEDICINE

[Pr]TYRUKO (pronounced tie-ROO-koh)

Natalizumab for injection

TYRUKO is a biosimilar biologic drug (biosimilar) to the reference biologic drug TYSABRI®. A biosimilar is authorized based on its similarity to a reference biologic drug that was already authorized for sale.

Read this carefully before you start taking TYRUKO and each time you get a TYRUKO IV infusion. This leaflet is a summary and will not tell you everything about this drug. Talk to your healthcare professional about your medical condition and treatment and ask if there is any new information about TYRUKO.

Keep this leaflet and the Patient Wallet Card. You should read them before starting TYRUKO, and before each TYRUKO IV infusion.

- It is important that you keep the Card with you during treatment and for three months after the last dose of TYRUKO, since side effects may occur even after you have stopped treatment.
- Show your Card and this package leaflet to any doctor involved in your treatment.

Serious Warnings and Precautions

- There have been uncommon cases of a brain infection by John Cunningham virus (JC virus) resulting in progressive multifocal leukoencephalopathy (PML) and/or JCV granule cell neuronopathy (JCV GCN) occurring in patients who have been given TYRUKO. These infections are associated with an uncontrolled increase of the JC virus in the brain, although reason for this increase in some patients treated with TYRUKO is unknown. It usually happens in people with weakened immune systems, but it is difficult to predict who will get these infections. Such infections may lead to severe disability or death; there is no known cure.
- In order to receive TYRUKO you must talk to your doctor and understand the benefits and risks of TYRUKO and consent to treatment prior to receiving your first treatment. After 24 months of treatment you should again talk to your doctor, understand the benefits and risks of TYRUKO treatment and consent to continuation of treatment.
- TYRUKO can only be given to patients who are registered in, and meet all conditions of the Sandoz PLUS Support Program. Sandoz PLUS Support Program is a controlled distribution program for TYRUKO.
- You should agree to enrol into the Sandoz PLUS Support Program, which is a patient registry, by contacting 1-888-449-7673.

What is TYRUKO used for?

TYRUKO decreases the number of MS attacks and slows down the progression of disabling effects of MS. Therefore, when you receive TYRUKO, you might not notice anything happening to your MS, but it may help to prevent your MS from becoming worse.

How does TYRUKO work?

TYRUKO is a man-made protein. It prevents the active immune cells from reaching the brain. TYRUKO is used for decreasing the inflammation in your brain (as seen on magnetic resonance imaging [MRI] scan) and therefore reduces nerve damage caused by multiple sclerosis.

What are the ingredients in TYRUKO?

Medicinal ingredients: natalizumab

Non-medicinal ingredients: L-histidine, L-histidine hydrochloride monohydrate, polysorbate 80, sodium chloride, water for injection.

TYRUKO comes in the following dosage forms:

Solution for intravenous administration 300mg/15mL.

TYRUKO comes in the form of a liquid in a vial. The liquid contains 300 mg in a 15 mL dose (20 mg/mL) of natalizumab. The liquid must be mixed with 0.9% sodium chloride and is administered into a vein over time, which is called infusion.

Do not use TYRUKO if:

- You have an allergy or are sensitive to natalizumab or anything else that is in this medicine (see Allergic reaction below).
- You have a serious problem with your immune system (for example, due to a disease such as leukemia or human immunodeficiency virus [HIV], or from using some other medicines that weaken your immune system).
- You have a serious infection, including an uncommon infection of the brain called progressive multifocal leukoencephalopathy (PML) now or in the past.
- You have active cancer.

To help avoid side effects and ensure proper use, talk to your healthcare professional before you take TYRUKO. Talk about any health conditions or problems you may have, including if you have:

- **Allergic reaction**
  Some patients have had allergic reactions to TYRUKO. If you notice any of the following signs of allergy to TYRUKO during or shortly after your infusion, tell your healthcare professional (doctor or nurse) immediately:
  - Itchy rash (hives)
  - Swelling of your face, lips or tongue
  - Difficulty breathing
  - Chest pain or discomfort
- **Infections**
  There have been uncommon cases of a brain infection by JC virus resulting in progressive multifocal leukoencephalopathy (PML) occurring in patients who have been given TYRUKO. PML is a serious condition, which may lead to disability or death. A condition called granule cell neuronopathy (GCN) is also caused by JC virus and has occurred in some patients who have been given TYRUKO. The symptoms of JCV GCN are similar to PML.

Your chance of getting PML increases:

- if you have antibodies against the JC virus, the virus that can cause PML. JC virus is a common virus which infects many people but does not normally cause noticeable illness. It is also very common to have these antibodies against the JC virus. If you do not have antibodies against the JC virus, you are at a lower risk of getting PML. Your doctor may recommend a blood test to see if you have these antibodies before you start TYRUKO. If you do not have the antibodies your doctor may repeat the test every 6 months while you are taking TYRUKO.
- with a longer period of TYRUKO treatment, especially if you have been on treatment for over 24 months.
- if you have received medicines that can weaken or suppress your immune system prior to starting TYRUKO (immunosuppressants), for example: azathioprine, cyclophosphamide, methotrexate, mitoxantrone, mycophenolate.

You must carefully consider and discuss with your physician the benefits and risks of TYRUKO therapy if you have ALL of the following risk factors: anti-JCV antibody positive, have received more than 2 years of TYRUKO therapy, AND have received medicines that can weaken or suppress your immune system (immunosuppressant therapy).

To monitor the risk of PML, your doctor may repeat the test regularly (eg. 6 months) to check if anything has changed if:

- You do not have antibodies to the JC virus in your blood

OR

- You have been treated for more than 2 years and you have a lower level of JCV antibodies in your blood.

A variety of symptoms of PML can appear and these can get worse over time. This is why it is important that you speak with your partner or caregivers and inform them about your treatment.

The symptoms of PML may be similar to an MS attack, including increasing weakness or clumsiness on one side of the body, trouble with vision, or trouble with thinking.

Therefore, if you feel your MS is getting worse, or if you notice any new symptoms, you should speak to your doctor immediately. Symptoms might arise that you might not be aware of yourself and may include changes in mood or behaviour, memory problems, speech and language difficulties, changes in your balance or walking ability. If any of these symptoms occur, it is important that you, your partner or caregiver inform your doctor as soon as possible. Based on this information your doctor may request further testing to rule out PML.

You and your caregiver should continue to watch for any signs and symptoms of PML for at least 6 months after you stop taking TYRUKO. Tell your doctor as soon as possible if you start noticing any symptoms.

It is not known if the chance of getting PML continues to rise, remains the same, or falls after you have been on TYRUKO for more than three years.

In most natalizumab treated patients with PML a reaction known as IRIS (Immune Reconstitution Inflammatory Syndrome) has occurred after stopping or removing natalizumab from the blood by a treatment called plasma exchange. IRIS presents as a worsening of your neurological symptoms that may be rapid and require that your doctor treat this condition with other medicines. IRIS can lead to serious

complications and may be fatal.

Because TYRUKO can weaken your immune system, you may have an increased chance of getting an unusual, serious or opportunistic infection (infection that usually does not cause disease in healthy people), such as herpes encephalitis and meningitis (inflammation of the brain and spinal cord). These infections can sometimes be life-threatening or fatal. Herpes infections of the eye have also occurred. Call your doctor right away if you have changes in vision, redness, or eye pain.

- **Liver or kidney problems**
  If you have problems with your kidneys, be sure to tell your doctor. If you experience unusual darkening of the urine, nausea, vomiting, feeling tired or weak and yellowing of the skin and eyes (jaundice), call your doctor right away.
- **Pregnancy**
  It is not known if TYRUKO can harm your baby if you are pregnant. You should not take TYRUKO if you are pregnant. Talk to your doctor if you become pregnant while taking TYRUKO.
- **Breastfeeding**
  TYRUKO has been found in breast milk. You should not breastfeed while taking TYRUKO. You should discuss with your doctor whether you should choose to breastfeed or to use TYRUKO.
- **Other considerations**
  TYRUKO is not intended for use in patients under the age of 18. TYRUKO has not been well studied in patients over 65 years old.
- Talk to your doctor if you are taking or have recently taken any other medications, including over-the-counter medicines or herbal (natural healthcare) products.
- TYRUKO can have an effect on the results of some laboratory tests showing an increase in the number of some blood cells.
- Talk to your doctor if you have easy bruising, unusual or prolonged bleeding from cuts, pinpoint or round spots that appear on your skin, abnormally heavy menstrual periods, or bleeding from the nose or gums that is new.
- Before you get TYRUKO IV, it is mixed with 0.9% sodium chloride. After mixing, each dose of TYRUKO contains 406 mg of sodium. This should be taken into consideration if you are on a controlled sodium diet.
- **Driving and using machines**
  TYRUKO is not expected to have an effect on your ability to drive or to operate machines. However, if you experience dizziness while taking TYRUKO, avoid driving or operating machines until it has resolved.

Tell your doctor about all of the medicines you take now or have taken in the last while, including those that are prescribed for you as well as those that you buy over-the-counter. It is not known if TYRUKO interacts with food or herbal (natural healthcare) products.

You may not be able to take TYRUKO with some medicines that affect your immune system.

Tell your healthcare professional about all the medicines you take, including any drugs, vitamins, minerals, natural supplements or alternative medicines.

The following may interact with TYRUKO:

Interactions with food and other drugs have not been established.

How to take TYRUKO IV:

TYRUKO can only be prescribed by a healthcare professional who is trained in treating neurological conditions. TYRUKO will be prepared and given to you by a healthcare professional.

Usual dose:

The usual adult dose is 300 mg given by intravenous infusion once every 4 weeks.

Overdose:

If you receive more TYRUKO than your healthcare professional prescribed, you should be monitored closely for any harmful signs or symptoms and given treatment for these right away, should they appear.

If you think you, or a person you are caring for, have taken too much TYRUKO, contact your healthcare professional, hospital emergency department, regional poison control centre or Health Canada's toll-free number, 1-844 POISON-X (1-844-764-7669) immediately, even if there are no signs or symptoms.

Missed Dose:

If you miss your usual dose of TYRUKO, contact your doctor to schedule your appointment as soon as possible. You should then continue to receive your dose of TYRUKO every 4 weeks.

What are possible side effects from using TYRUKO?

These are not all the possible side effects you may have when taking TYRUKO. If you experience any side effects not listed here, tell your healthcare professional. Show your Wallet Card and this package leaflet to any doctor involved in your treatment.

Serious side effects and what to do about them			
Symptom / effect	**Talk to your healthcare professional**		**Stop taking drug and get immediate medical help**
	Only if severe	**In all cases**	
VERY COMMON			
Dizziness	✓		
Feeling sick (nausea)	✓		
Headache	✓		
Joint pain	✓		
Sore throat and runny or blocked up nose	✓		
Tiredness	✓		

Serious side effects and what to do about them			
Symptom / effect	**Talk to your healthcare professional**		**Stop taking drug and get immediate medical help**
	Only if severe	**In all cases**	
Urinary (bladder) infection		✓	
COMMON			
Being sick (vomiting)	✓		
Fever		✓	
Itchy rash (hives)		✓	
Shivering		✓	
UNCOMMON			
Progressive multifocal leukoencephalopathy (PML), a rare brain infection. Typical symptoms include: • progressive weakness on one side of the body • clumsiness of limbs • disturbance of vision • changes in thinking, memory and orientation • confusion • personality changes			✓
Reduction in blood platelets and easy bruising (purpura)		✓	
Severe allergy (hypersensitivity)			✓
RARE			
Liver symptoms		✓	
Severe anemia (decrease in red blood cells). Symptoms include pale skin, feeling breathless, lack of energy		✓	
Unusual infections			✓

If any of these occur during or shortly after the infusion, tell your doctor or nurse immediately.

Some patients have had allergic reactions during or shortly after receiving natalizumab. Your doctor or nurse will stop your TYRUKO infusion if he or she sees any signs or symptoms of an allergic reaction.

After you have received TYRUKO, a doctor or nurse may monitor you for 1 hour. Speak to your doctor as soon as possible if you think you have an infection.

If you have a troublesome symptom or side effect that is not listed here or becomes bad enough to interfere with your daily activities, tell your healthcare professional.

Reporting Side Effects

You can report any suspected side effects associated with the use of health products to Health Canada by:

- Visiting the Web page on Adverse Reaction Reporting (canada.ca/drug-device-reporting) for information on how to report online, by mail or by fax; or
- Calling toll-free at 1-866-234-2345.

NOTE: Contact your health professional if you need information about how to manage your side effects. The Canada Vigilance Program does not provide medical advice.

Storage:

Unopened vial: Store in a refrigerator at 2°C to 8°C. Do not freeze. Keep the vial in the outer carton to protect it from light. Do not shake. Do not use after the expiry date found on the label and carton.

Diluted solution: After your healthcare professional has prepared TYRUKO for infusion, the diluted solution must either be used immediately or should be stored in a refrigerator (2°C to 8°C). Infusion of the diluted product should be started as soon as possible and completed within 24 hours of dilution.

Keep out of reach and sight of children.

If you want more information about TYRUKO:

- Talk to your healthcare professional
- Find the full product monograph that is prepared for healthcare professionals and includes this Patient Medication Information by visiting the Health Canada Drug Product Database website (Drug Product Database: Access the database); the manufacturer's website (www.sandoz.ca), or by calling Sandoz Canada Inc. (1-800-361-3062) or Sandoz PLUS (1-888-449-7673).

This leaflet was prepared by Sandoz Canada Inc.

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